MWNR

MORBIDITY AND MORTALITY WEEKLY REPORT

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Wound Botulism — California, 1995

During January–November 1995, a total of 19 laboratory-confirmed cases of wound botulism were reported to the California Department of Health Services (CDHS); of these, 13 had occurred since August. Since 1990, the number of wound botulism cases reported annually in California has increased steadily (one case in 1990, two in 1991, three in 1992, four in 1993, and 11 in 1994). All cases except one since 1991 have occurred in injecting-drug users, and many involved subcutaneous injection or "skin popping" of black tar heroin. This report summarizes the findings of the investigation of two cases.

Case 1

On September 23, a 44-year-old male user of black tar heroin developed an abscess on his right arm, which was treated unsuccessfully with cephelexin and ciprofloxacin; on September 29, the abscess was incised and drained. On October 1, he was examined at a local emergency department (ED) because of slurred speech and was released.

On October 3, he sought care in the ED of a community hospital in Yolo County because of difficulty swallowing, which progressed to slurred speech, blurred vision, neck and arm weakness, and shortness of breath. Findings on physical examination included ophthalmoplegia; ptosis; and weakness of his facial, sternocleidomastoid, and deltoid muscles. Examination of a sample of his cerebrospinal fluid detected a marginally elevated protein level (50 mg/dL). A "Tensilon®* test" (intravenous administration of edrophonium bromide to improve strength) was negative, and electromyography was not performed. Despite treatment with intravenous gamma globulin for suspected Guillain-Barré syndrome, weakness progressed, and on October 4, he required mechanical ventilation. On October 5, the diagnosis of wound botulism was considered, and CDHS was consulted. Two vials of botulinal antitoxin were released by CDHS and administered to the patient; in addition, treatment with 12 million units of penicillin daily was initiated.

^{*}Use of trade names and commercial sources is for identification only and does not imply endorsement by the Public Health Service or the U.S. Department of Health and Human Services.

Wound Botulism - Continued

A serum specimen obtained from the patient on October 4 was positive for type A botulinal toxin by mouse bioassay. No tissue from the abscess could be obtained for culture. The patient was discharged on November 21.

Case 2

On September 25, a 30-year-old pregnant woman who reported last skin popping black tar heroin on September 24 sought care at an ED in Ventura County because of a sore throat and the sensation of "heavy eyelids." An upper respiratory tract infection was diagnosed, and she was released. On September 27, she developed difficulty swallowing and speaking and was admitted to a community hospital for evaluation. During the 12 hours following admission, she developed ophthalmoplegia and profound, symmetric, p. oximal paralysis of arms and legs, affecting her arms more than her legs; she subsequently required mechanical ventilation. A Tensilon® test was negative. Electromyography with repetitive motor-nerve stimulation at 10 Hz increased the muscle action potential by 17%. Lumbar puncture could not be performed.

On September 29, she underwent wide excision of multiple abscesses on her left leg. Botulism was suspected; CDHS was consulted and released two vials of antitoxin for administration to the patient. Treatment with high-dose penicillin was initiated.

Tissue and serum specimens obtained from the patient were positive for type A botulinal toxin by mouse bioassay, and histochemical staining of an excised abscess indicated the presence of spores and vegetative cells consistent with *Clostridium botulinum*. Culture of tissue from the wound yielded *C. botulinum* type A. On November 21, the patient was discharged from the hospital; her baby, who was delivered by cesarean section at 34 weeks on November 11, remained in intensive care on December 7.

Reported by: M Gollober, MD, RA Beyer, MD, Woodland Memorial Hospital, S Kwan, RO Bates, MD, Yolo County Health Dept, Woodland; H Oster, MD, Community Memorial Hospital, M Billimek, SE Matson, G Feldman, MD, Ventura County Health Dept, Ventura; R Bryant, J McGee, SB Werner, MD, CA Glaser, MD, DJ Vugia, MD, SH Waterman, MD, State Epidemiologist, California Dept of Health Svcs. Div of Field Epidemiology, Epidemiology Program Office, CDC. Editorial Note: Wound botulism, first described in association with traumatic injury, is a rare illness that occurs after spores of C. botulinum have germinated in a wound and produced botulinal toxin, resulting in flaccid paralysis (1). Wound botulism attributable to drug injection was first reported in 1982 in New York City (2); since then, such cases have been reported only sporadically. However, wound botulism occurred in 11 (21%) of the 53 botulism cases among adults reported to CDC in the United States in 1994, and all occurred among injecting-drug users in California.

Black tar heroin is dark and gummy. The drug available in California is believed to be processed in facilities close to the source of opium poppies grown in several states in Mexico. The final product often contains adulterants as well as diluents (e.g., sugar) to increase bulk. The use of black tar heroin is believed to be increasing and, since 1993, has supplanted traditional forms of heroin in California and other western states. However, it is unknown whether the increase in cases of wound botulism reflects increased supply of the drug, a change in its manufacture and distribution, or a change in drug-using behavior.

Skin popping of heroin is common among chronic users who are either unable or reluctant to inject the drug intravenously. Unlike botulinal toxin, which is inactivated by heat, spores of *C. botulinum*—which could be in the heroin or in the liquid (usually

Wound Botulism - Continued

water) with which the heroin is dissolved—are not destroyed by heating the heroin/liquid mixture. Spores inoculated into subcutaneous tissue—either from the drug or from the skin after inadequate skin disinfection—can germinate and produce toxin.

Botulism should be suspected in patients with acute onset of flaccid paralysis with ophthalmoplegia, ptosis, or other cranial nerve dysfunction, particularly when the paralysis is descending, symmetric, and associated with a normal cerebrospinal fluid protein level. A history of drug injection or a food history that does not identify a probable source for foodborne botulism should prompt consideration of wound botulism and elicitation of a thorough history and physical examination for evidence of cellulitis or abscess. A meticulous physical examination is necessary because wounds containing *C. botulinum* may be small and initially unnoticed. Inspection of the intranasal septum and paranasal sinuses also may disclose a focus of *C. botulinum* infection in persons who snort cocaine (3). The diagnosis is supported by either conventional electromyography showing potentiation after supramaximal stimulation at 20–50 Hz, or single-fiber electromyography showing increased jitter and blocking (4). A diagnosis of myasthenia gravis would be supported by improvement in muscle function after the administration of edrophonium bromide (Tensilon®). Initial treatment decisions should not necessarily await neurologic test results.

Both risk for death and duration of hospitalization can be reduced by prompt administration of botulinal antitoxin (5). The administration of antitoxin is not contraindicated by pregnancy. Wounds suspected of being contaminated with *C. botulinum* should be widely debrided and irrigated, ideally after the administration of botulinal antitoxin. Penicillin, 10–20 million units per day, is considered the antibiotic of choice, although its efficacy has not been determined (6). Mechanical ventilation is the main supportive therapy for treatment of severe botulism.

Because of the increase in wound botulism cases, CDHS has publicized this problem through press releases and provided informational materials for county health officials, ED physicians, and community-based organizations offering outreach to drug users. Clinically suspected cases of botulism should be reported *immediately* to local or state public health agencies to facilitate 1) laboratory confirmation of the diagnosis (using serum and tissue specimens for suspected wound botulism; stool and possibly serum specimens for suspected infant botulism; and food, serum, stool, and gastric aspirate specimens for suspected foodborne botulism); 2) release of antitoxin, if clinically indicated; and 3) prompt investigation of all likely foodborne sources to identify and eliminate a suspected food source to protect other persons. In addition, injecting-drug users should be reminded of the health risks associated with illicit drug use, including the possibility of botulism.

If local and state officials are not available, CDC can be contacted directly (telephone [404] 639-2206, Monday through Friday, 8 a.m.-4:30 p.m. Eastern Time or [404] 639-2888 at other times). In California, health-care workers should contact CDHS (telephone [510] 540-2308), where consultation is available at all times for suspected botulism cases.

References

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Wound Botulism - Continued

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Unexplained Severe Illness Possibly Associated with Consumption of Kombucha Tea — Iowa, 1995

Kombucha tea is a popular health beverage made by incubating the Kombucha mushroom in sweet black tea. Although advocates of Kombucha tea have attributed many therapeutic effects to the drink (1–3), its beneficial and/or adverse effects have not been determined scientifically. During April 1995, cases of unexplained severe illness (including one death) occurred in two persons in a rural town in northwestern lowa who had been drinking Kombucha tea daily for approximately 2 months. Based on the findings of a preliminary investigation by the lowa Department of Public Health (IDPH), on April 10 IDPH issued a news release recommending that persons refrain from drinking Kombucha tea until the role of the tea in the two cases of illness had been evaluated fully. This report summarizes the investigation of these cases by the IDPH, CDC, and the Food and Drug Administration (FDA).

Patient 1

On April 1, a 59-year-old woman was found unconscious in her home by a neighbor and was transported to a local hospital. On arrival in the emergency department, respiratory therapy was initiated with oxygen. Her family members reported that, 1 hour earlier, she appeared fatigued but had no specific medical complaints. Analysis of arterial blood samples indicated severe metabolic acidosis; her pH level was 6.9 (normal: 7.37–7.43); pO₂, 474.9 mm Hg (normal: 75–80 mm Hg); and pCO₂, 39.2 mm Hg (normal: 35–45 mm Hg). She also had elevated levels of lactic acid (9.85 mM [normal: 0.67 mM–2.47 mM]) and a base excess of –19.5 (normal: –2—+2). Her daughter and her primary physician reported that she took medications for hypertension, anemia, and mild renal insufficiency. Soon after admission, symptoms of disseminated intravascular coagulopathy began; she suffered cardiac arrest and was resuscitated, but her condition continued to deteriorate. She died on April 3.

The cause of the woman's acute metabolic disorder was not established. An autopsy detected evidence of peritonitis with fecal contamination of the peritoneal cavity, although the location of perforation could not be determined. Neither the woman's clinical history nor autopsy findings supported a cardiogenic cause. Toxicologic analyses for a series of prescription and nonprescription drugs and carbon monoxide and cyanide poisoning were negative. Her daughter reported that, during the previous 2 months, the patient had drank approximately 4 oz of Kombucha tea daily.

Kombucha Tea — Continued

Patient 2

On April 10, a previously healthy 48-year-old woman had onset of shortness of breath and was transported by ambulance to the same hospital as patient 1. On admission, she was in respiratory distress. Chest radiographs revealed extensive acute pulmonary edema. Analysis of arterial blood samples indicated severe metabolic acidosis with uncompensated respiratory acidosis; her pH level was 6.7; pO₂, 86 mm Hg; and pCO₂, 67 mm Hg. She had elevated levels of lactic acid (12.4 mM) and a base excess of –28. The woman suffered cardiac arrest but was resuscitated and stabilized. She improved and was discharged on April 13.

Toxicologic analyses for a series of prescription and nonprescription drugs were negative, and there was no evidence of a septic or cardiogenic cause. The patient reported drinking Kombucha tea during the previous 2 months and had obtained her original mushroom from the same person as patient 1. On April 10, immediately before the onset of illness, she had increased the amount of tea she consumed from 4 oz daily to 12 oz, and she had increased the period of incubation for that batch of tea from 7 days to 14 days.

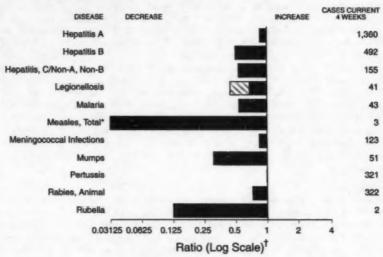
Investigation

The mushrooms used by both women were derived from the same parent mushroom. At least 115 additional persons in the town had used or were using mushrooms from the same source as for the two ill women, but no other cases of unexplained acute illness were reported among these persons. A review of hospital emergency department records for March 1–April 10 did not detect other cases of unexplained lactic acidosis or other likely cases of tea-associated acute illness.

Samples of the mushrooms and samples of the tea consumed by both casepatients were sent to FDA for analysis. Microbiologic analysis of the tea and mushrooms identified several species of yeast and bacteria, including Saccharomyces cerevisiae and Candida valida. No known human pathogens or toxin-producing organisms were identified. The alcohol content of the tea ranged from 0.7% to 1.3%; no methanol was detected.

To characterize the methods used for preparing the tea, IDPH and CDC surveyed a nonrandom sample of 24 persons in the town who regularly drank Kombucha tea. The average age of survey participants was 57.1 years. Of the 21 participants for whom information was available, 20 had obtained their mushrooms from friends or relatives, and 15 (71%) of these had given mushrooms to their friends. One person had purchased a mushroom from a commercial producer. Of the 20 participants who had prepared the tea themselves, most (12 [60%]) reported incubating the Kombucha mushroom at room temperature for 7–10 days in 3 quarts of sweetened tea and drinking 4 oz of it per day. Patient 1 followed this regimen; patient 2 had incubated the mushroom longer (14 days) and consumed more tea (12 oz per day). Five (25%) other persons who had prepared their own tea reported incubating the mushroom for 13–14 days, and two (8%) of the 24 total participants reported consuming up to 8 oz of tea per day. Of the 21 persons for whom information was available, five (23%) discarded batches of tea because of their concerns about the appearance or taste of the tea or because of visible mold growth.

FIGURE I. Notifiable disease reports, comparison of 4-week totals ending December 2, 1995, with historical data - United States



Beyond Historical Limits

*The large apparent decrease in the number of reported cases of measles (total) reflects dramatic fluctuations in the historical baseline.

[†]Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary — cases of specified notifiable diseases, United States, cumulative, week ending December 2, 1995 (48th Week)

	Cum. 1995		Cum. 1995
Anthrax Brucellosis Cholers Congenital rubella syndrome Diphtheria Haemophilus Influenzae* Hansen Disease Piaque	78 16 6 1,061 125	Paittacosis Rabies, human Rocky Mountain Spotted Fever Syphilia, congenital, age < 1 year ¹ Tetanus Toxic shock syndrome Trichinosis Typhoid fever	64 2 543 469 30 165 28 305

*Of 1,038 cases of known age, 250 (24%) were reported among children less than 5 years of age.

*Updated quarterly from reports to the Division of STD Prevention, National Center for Prevention Services. This total through
third quarter 1995.

-: no reported cases

TABLE II. Cases of selected notifiable diseases, United States, weeks ending December 2, 1995, and December 3, 1994 (48th Week)

Reporting Area	AIDS*	Gonor	rhea	A				C/NA	,NB	Legionellosis		
	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1984	Cum. 1996	Cum. 1994	Cum. 1996	Cum. 1994	Cum. 1995	Cum. 1994	
UNITED STATES	65,706	321,463	373,705	26,529	23,067	8,956	10,602	3,379	3,758	1,088	1,454	
NEW ENGLAND	3,119	5,783	7,808	297	272	190	316	12	138	35	74	
Maine	81	80	87	30	24	12	11			6	5	
N.H. Vt.	87 30	104	101	11	16 12	20	25 11	12	10 15	2	i	
Mass.	1,339	2,677	3,012	129	100	83	174		93	22	62	
R.I.	214	501	443	34	25	8	8		20	5	16	
Conn.	1,368	2,361	3,931	88	95	66	87	-	*	N	N	
MID. ATLANTIC	17,668	32,842	42,243	1,599	1,553	1,181	1,436	427	429	178	243	
Upstate N.Y. N.Y. City	2,127 9,225	3,853 11,818	10,599 15,006	432 737	506 603	357 349	349	232	204	50	57	
N.J.	4,158	5,391	4,799	231	268	302	356	154	186	5 27	43	
Pa.	2,158	11,780	11,839	199	176	173	362	40	35	96	136	
E.N. CENTRAL	4,940	67,066	75,689	2,836	2,426	966	1,103	241	302	291	411	
Ohio	1,017	18,238	20,229	1,673	993	101	154	15	23	142	189	
ind.	499 2.054	7,612	8,484	170	344	234	198	2	9	71	45	
Mich.	1,039	19,281 16,796	22,764 16,943	479 343	570 296	202 373	287 374	160	78 192	16 32	40 76	
Wis.	331	5,139	7,269	171	223	58	90	100	192	30	61	
W.N. CENTRAL	1.555	17,620	20,722	1,719	1,122	552	620	140	89	108	98	
Minn.	347	2,609	3,244	173	224	62	59	4	17	6	3	
lowa	94	1,429	1,423	65	58	45	25	12	13	21	30	
Mo.	713	10,242	11,339	1,180	573	364	474	98	28	51	40	
N. Dak. S. Dak.	5 18	26 206	37 211	24 79	35	2	2	8	1	4	4	
Nebr.	101	757	1,060	46	119	31	28	6	13	14	14	
Kans.	277	2,351	3,408	152	108	44	31	11	17	8	6	
S. ATLANTIC	16,629	96,395	99,851	1,236	1,230	1,357	1,917	317	418	166	347	
Del.	279	2,079	1,872	8	22	8	14	-	2	2	31	
Md. D.C.	2,409 976	8,852 4,369	17,162 6,572	210	177 25	240	322 50	4	20	30	78	
Va.	1,400	9,601	12,424	194	174	103	124	18	25	18	13	
W. Va.	116	599	758	24	22	52	44	43	41	4	4	
N.C.	951	21,574	26,330	104	139	286	250	58	54	31	27	
S.C.	868 2,144	11,502	12,171	44	39 40	49	32	16	10	30	16	
Fla.	7,486	18,799 19,020	22,562	54 577	592	62 538	543 529	13 165	196	14 32	110	
E.S. CENTRAL	2.093	38,116	43,062	1,747	632	745	1,119	864	854	43	81	
Ky.	267	4,515	4,793	39	160	63	74	23	29	10	9	
Tenn.	843	12,571	14,243	1,435	290	579	964	839	807	24	43	
Ala.	562	15,351	13,604	80	109	103	81	2	18	6	13	
Miss.	421	5,679	10,422	193	73					3	16	
W.S. CENTRAL Ark.	5,626	31,074	45,229	4,263	2,925	1,343	1,209	306	296	18	41	
La.	972	3,445 9,881	6,191	597 140	140	71 202	24 153	139	167	3	13	
Okia.	256	4,955	4,475	1,068	355	206	124	73	55	6	11	
Теж.	4,155	12,793	23,471	2,458	2,249	864	908	90	67	8	9	
MOUNTAIN	2,071	7,567	9,464	3,830	4,671	739	614	370	426	107	90	
Mont.	22	65	84	163	23	22	19	13	13	4	16	
Idaho Wyo.	43 15	114	81 83	301 101	353 29	82 25	71 23	147	67 161	12	5	
Colo.	631	2,627	3,316	492	539	130	91	54	74	38	18	
N. Mex.	155	945	986	745	1.023	268	198	42	45	4	4	
Ariz.	635	2,848	3,047	1,201	1,897	102	79	46	30	12	14	
Utah Nev.	143 427	131 788	281 1,586	635 192	578 229	72 38	78 55	10 17	18	17 18	7 24	
PACIFIC	12,004	25,000	29,837	9,002	8,236	1,881	2,268	702	804		-	
Wash.	855	2,381	2,679	770	989	1,881	2,208	204	248	122	12	
Oreg.	426	364	942	2,116	1,046	107	143	31	41	-		
Calif.	10,441	20,817	24,729	5,916	5,941	1,555	1,872	463	510	96	53	
Alaska Hawaii	62 220	629 809	844 643	51 149	205 55	10 29	13 26	2 2	5	5	4	
	220					-	20	2	9			
Guarn P.R.	2,189	77 540	127 463	6 89	23 81	488	309	18	183	1	1	
V.I.	30	6	403	89	3	400	309	18	163	-		
Amer. Samoa	-	35	31	6	9	-						
C.N.M.I.		42	46	18	12	13	1		-			

N: Not notifiable U: Unavailable : no reported cases C.N.M.I.: Commonwealth of Northern Mariana Islands
*Updated monthly to the Division of HIV/AIDS Prevention, National Center for Prevention Services, last update November 30, 1995.

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending December 2, 1995, and December 3, 1994 (48th Week)

Reporting Area	lame.						Measi	es (Rube						
	Lyn	me iase	Malaria		Indigenous		Impo	orted*	To	tal	Infec	ococcai tions	Mumps	
	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	1995	Cum. 1995	1995	Cum. 1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum 1994
UNITED STATES	8,546	11,727	1,166	1,003		261		26	287	887	2,691	2,494	761	1,319
NEW ENGLAND	1,967	2,690	47	71	-	8	-	2	10	27	131	122	11	25
Maine	28	27	7	6						5	10	21	4	3
N.H. /t.	26	28 16	2	3	-					3	23	- 4	1	4
Mans.	191	198	18	33	-	2		1	3	7	43	57	2	3
R.I.	285	489	4	9		5			5	7			1	3
Conn.	1,431	1,952	15	17		1	-	1	2	4	44	32	3	12
MID. ATLANTIC	5,437	7,193	318	215	-	7		5	12	223	304	271	107	106
Upstate N.Y.	2,641	4,429	170	52 77	-	1 2		3	5	26 15	96	89	25	3
N.Y. City N.J.	1,303	1,417	62	50		4	-	2	6	173	76	32 55	15 13	10
Pa.	1,267	1,319	25	36	-					9	88	95	54	58
E.N. CENTRAL	87	523	131	99		13		4	17	102	388	364	165	231
Ohio	52	44	11	15		1	-	1	2	17	110	107	51	- 00
Ind.	20	18	17	13			*			1	51	48	10	7
III. Mich.	10	23 31	63 26	42 26		4 6		2	6	56 25	92 70	116	46 58	104
Wis.		407	14	3		2	-		2	3	45	56 37	56	46
W.N. CENTRAL	254	281	26	45		2			2	170	179	161	47	67
Minn.	174	150	6	14	-					170	27	20	8	
lowa	15	16	2	5			-			7	30	19	10	10
Mo.	40	99	8	13		1		*	1	160	73	75	23	40
N. Dak. S. Dak.			2 2	1	-						8	9	1	
Nebr.	3	3	3	5		-				2	15	13	4	
Kans.	22	13	3	7		1	*		1	ī	25	24	1	
S. ATLANTIC	518	784	230	217		11		1	12	72	501	367	98	19:
Del.	23	105	1	3		*					6	5		
Md.	286	294	60	78		*		1	1	4	34	32	20	6
D.C. Va.	53	127	16 52	14 36	*			*		3	7 59	66	25	4
W. Va.	23	26	4	30						37	8	12	25	-
N.C.	82	77	16	11		-				3	80	51	16	3
S.C.	17	. 7	3	5		-					57	31	11	
Ga. Fla.	14	119		33 37	-	2 9	-		9	21	102 148	76 88	10	3
E.S. CENTRAL	50	43		31										
Ky.	10	24		11			-		:	28	168 53	179 36	20	2
Tenn.	24	13		10						28	41	35	5	
Ala.	9	6	9	9							41	73	4	1
Miss.	7	-	3	1						*	33	35	11	
W.S. CENTRAL	111	122		42		31	**	3		19	327	301	53	22
Ark. La.	9 7	8 2		3	-	17		i	18	1	31 49	43 39	10	3
Okla.	48	73		7		17			10		38	33	13	2
Tiese.	47	39				12		2	14	17	209	186	30	16
MOUNTAIN	12	17	58	35		68	-	2	70	165	184	167	25	15
Mont.			. 3								4	6	1	
Idaho		3			-	1		1		1	11	17	3	1
Wyo. Colo.	3	5		15		26			26	19	7 45	38	2	
N. Mex.	i	5				30	-	1		10	35	15	Ñ	
Ariz.	1		12	8		10	-		10	2	58	55	2	
Utah	1	2				i			:	134	15	19	11	3
Nev.	5			_			-		1	9	9	10	6	1
PACIFIC Wash.	110					121	-	9		81	529	562		27
Oreg.	13			16		10	-	1		2	96 98	130	14 N	1
Calif.	87	84	228			105		3		61	330		198	23
Alaska			. 3				*			10	12	3	13	
Hawaii			- 10	14			*	1	1 1	4	4			1
Guam			. :	:	U		U			228	3		4	
P.R. V.I.			- 1	5	Ü	11	Ú		- 11	11	23	7	2 2	
Amer. Samoa					ŭ		Ü							
C.N.M.I.			. 1	1	ŭ		Ü			29				

^{*}For imported measles, cases include only those resulting from importation from other countries.

N: Not notifiable U: Unavailable -: no reported cases

TABLE II. (Cont'd.) Cases of selected notifiable diseases, United States, weeks ending December 2, 1995, and December 3, 1994 (48th Week)

Reporting Area		Portuosis			Rutiella		Sypi (Prim Secon	hillis ary & idary)	Tubero	ulosis	Rabies, Animal		
	1995	Cum. 1995	Cum. 1994	1995	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	Cum. 1995	Cum. 1994	
UNITED STATES	64	3,869	3,906	-	139	215	13.603	19,290	18,370	20,270	6,288	7,185	
NEW ENGLAND	29	552	613		49	131	242	207	465	465	1,407	1,770	
Maine	1	46	18		1		2	4	12	27	45	1,770	
N.H.	7	53	82		1	*	1	4	18	14	143	199	
/t. Mass.	21	64	46		-	***	.:	-	4	8	170	137	
R.I.	21	358	422		7	125	64	87 15	259 45	240	393	687	
Conn.		27	39		40	3	171	97	127	132	311	707	
MID. ATLANTIC	1	357	611		14	7	782	1,305	3,743	4,174	1,200		
Jpstate N.Y.		200	227		5	6	44	162	477	561	474	1,925	
N.Y. City		33	176		8	-	368	562	1,974	2,398	-	1,10	
N.J.		14	15		1	1	163	220	722	739	316	261	
-	1	110	193		-		207	361	570	476	410	230	
E.N. CENTRAL	4	461	546		5	9	2,333	2,843	1,884	1,921	91	66	
Ohio ind.	1	153 73	146 63	-	1	*	787	1,069	256	306	12	4	
II.	2	113	101		1	1	267 834	237 980	305 892	170 968	13	13	
Mich.	1	110	92		3	8	287	278	362	424	40	13	
Wis.	-	12	144				158	279	69	53	11	15	
W.N. CENTRAL	1	247	199	-	1	2	678	1,102	521	519	328	215	
Minn.	-	127	87			-	36	45	124	122	24	19	
lowa		12	21				43	65	57	57	119	83	
Mo.	*	53	42			2	562	926	206	223	23	2	
N. Dak. S. Dak.		12	5 22	-			-	1	5	9	28	13	
Nebr.	1	12	9			-	11	11	22	24	86	3	
Kans.	-	23	13		1		26	52	21 86	17 67	5 43	34	
S. ATLANTIC		319	335		25	16	3,477	5,021	2,976				
Del.		10	3		20	10	16	25	46	3,487	1,977	1,872	
Md.	-	38	68				196	298	268	317	299	49	
D.C.		6	10	-			97	199	96	103	11		
Va.	*	31	36	-			550	751	255	292	418	40	
W. Va. N.C.	*	110	5 79		i	-	10	9	66	74	110	7.	
S.C.		27	14	1	i		1,050 541	1,536 745	416 294	461 355	435 118	150	
Ge.		29	30			2	661	764	319	618	264	34	
Fla.		68	90		23	14	356	694	1,216	1,207	238	150	
E.S. CENTRAL	1	270	128				3,436	3,661	1,449	1,476	274	21	
Ky.		24	60				187	195	286	292	28	2	
Tenn.		207	22	-	-		817	976	372	519	92	7	
Ala. Miss.	1	36	34 12	Ñ	N		606	613	378	400	145	114	
				14		N	1,826	1,877	413	265	9	4	
W.S. CENTRAL Ark.		280 41	185 27		8	13	1,872	4,102	2,576	2,742	521	64	
La.		17	10		1		97 962	439 1,577	208 105	233 193	43	3	
Okla.		31	27			4	181	148	330	222	28	3	
Tex.		191	121		7	9	632	1,938	1,933	2,094	450	50	
MOUNTAIN	7	529	491		5	5	202	229	594	534	162	14	
Mont.		9	10	-			4	3	10	9	43	2	
Idaho	1	96	80		-	*		1	15	11	3		
Wyo.	-	1	240		1		1	2	4	8	25	1	
Colo. N. Mex.	5	103 139	219 32				98 31	116 21	66 72	92 66	9	1	
Ariz.	-	149	112		3		35	45	300	201	50	5	
Uteh .		27	35		1	4	4	11	37	41	15	1	
Nev.	-	5	3	-		1	29	30	90	106	11		
PACIFIC	21	854	798		32	32	581	820	4.162	4,972	328	33	
Wash.	21	321	106	-	2		15	32	220	239	7	1	
Oreg.		59	102	-	2	4	9	35	68	90		1	
Calif. Alaska		415	571	*	24	24	556	746	3,648	4,338	317	26	
Hawsii	-	58	19		4	4	1	3 4	63 165	84 221	4	3	
Guam	U	1	2	- 11									
P.R.	1	15	3	U		1	288	3 295	53 195	75 189	47	7	
V.I.	Ü		-	Ü	-		200	28	195	109	4/	,	
Amer. Samoa	U		1	U				1	5	4			
C.N.M.I.	U	-		U	-		12	2	16	30			

U: Unavailable -: no reported cases

TABLE III. Deaths in 121 U.S. cities,* week ending December 2, 1995 (48th Week)

	A	il Cau	see, By	Age (Y	ears)		P&f	M*	All Causes, By Age (Years)						
Reporting Area	All 285 45-64		25-44	1-24	<1	Total	Reporting Area	All Ages	2:05	45-G4	25-44	1-24	<1	P&d Tota	
NEW ENGLAND loston, Mess. Irridgeport, Conn. Jambridge, Mess. Sill River, Mess. Sartford, Conn. Jowell, Mess. Lynn, Mess. Lynn, Mess. Lynn, Mess. Jower Haven, Conn. Providence, R.I. Somerville, Mess. Springfield, Mess. Springfield, Mess.	42 73 5 63 37	474 75 37 15 30 45 20 15 27 23 51 34 46 29	125 21 12 7 3 14 5 1 3 12 15 2 15 4	64 19 3 1 1 11 2 7 5	13 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	961	51 9 2 2 4 4 2 2 2 2 2 1 9 1 4 3	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Patersburg, Fla. Tampa, Fla. Washington, D.C. Wilmington, Del.	1,163 150 208 78 170 118 48 86 40 62 195 U	787 88 145 60 110 79 30 59 17 42 147 U	212 37 29 14 41 21 9 14 9 11 27 U	116 18 30 3 16 8 3 9 11 2 16 U	28 3 3 1 2 7 3 3 1 3 2 0 0	20 4 1 1 3 1 2 4 3 U	1
Norceeter, Mass. MID. ATLANTIC Albany, N.Y. Allientown, Pa. Buffalo, N.Y. Carnden, N.J. Elizabeth, N.J. Erie, Pa.\$	78 2,926 55 20 100 48 29 71	1,968 36 16 79 25 17 58	528 13 3 14 12 9 12	7 322 4 1 4 4 4 3	1 2 1	48 2	8 152 2 5 4 1 4	E.S. CENTRAL Birmingham, Ala. Chattanooga, Tenn. Knoxville, Tenn. Lexington, Ky. Memphia, Tenn. Mobile, Ala. Montgomery, Ala. Nashville, Tenn.	832 134 77 105 77 140 87 49 163	555 89 50 73 44 95 61 32	192 28 17 23 27 28 18 12 39	57 11 6 7 2 12 6 3 10	13 1 3 1 1 5 5	15 5 1 1 3 . 2 2 1	1 1
Jersey City, N.J. Hene York City, N.Y. Hener Kork City, N.Y. Henerak, N.J. Heterson, N.J. Hisbourgh, Pa.S Heading, Pa. Rochester, N.Y. Scranton, Pa.S Hyracuse, N.Y. Trenton, N.J. Lica, N.Y. Yonkers, N.Y.	78 1,489 50 32 400 92 15 184 21 25 124 49 14	48 961 14 21 284 65 11 149 18 17 96 35 13	1 6 15 7	7 202 18 3 37 5 3 9 2 2 9 6	1 36 2 1 8 3 3	2 21 3 3 2 2 1 1	3 58 3 6 28 9 2 12 1 1 10 2	W.S. CENTRAL Austin, Tex. Beton Rouge, La. Corpus Christi, Tex. Dellas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, Le. Sen Antonio, Tex. Shreveport, La. Tulsa, Okla.	1,708 96 86 47 233 65 131 385 86 101 268 76 134	1,045 65 48 37 137 37 89 223 43 52 189 50 95		180 11 14 1 29 5 16 36 12 16 26 5	58 2 3 13 3 5 14 4 2 11	43 2 4 5 7 9 6 2 6 1	36
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, Ill. Cincinnati, Ohio Cevesland, Ohio Columbus, Ohio Deyton, Ohio Deyton, Ohio Deyton, Ohio Deyton, Ohio Ceransville, Ind. Fort Wayne, Ind. Grand Rapids, Mic Indianapolis, Ind.	255	56 28 254 122 138 150 117 162 39 33 17 57	12 5 81 45 49 48 32 57 8 1 12 8 9	4 1 4 21	72 1 13 5 1 5 8 12 1 1 4	56 1 1 22 3 2 7 2 5 1	7 2 1 2 9	MDUNTAIN Albuquerque, N.M. Colo. Springe, Colo Denver, Colo. Las Veges, Nev. Ogden, Utah Phoenic, Ariz. Pueblo, Colo. Salt Lake City, Utah Tucson, Ariz. PACIFIC Berkeley, Calif. Fresno, Calif. Glendels, Calif.	99 211 37 218 34	1,037 1,037 14 74	16 8 14 52 49 6 23 22 266 5 13 U	109 19 8 16 15 4 25 1 13 8 132 121 U	25 3 2 3 2 2 6 7	24 3 2 5 2 7 4 1 1 29 1 3 0 0	1
Madiason, Wis. Miliwaukee, Wis. Peoria, III. Peoria, III. South Bend, Ind. Toledo, Ohio W.N. CENTRAL Des Moines, Iowa Des Moines, Iowa Duiuth, Minn. Kansas City, Kansas City, Mo. Lincoln, Nebr. Minnespolis, Minn Omaha, Nebr. St. Louis, Mo.	712 63 26 26 20 83	1 1256 306 306 306 306 306 306 306 306 306 30	288 188	15 3 4 5 1 5 1 8 1 2 9 9	1 4	1	2 6 5 7 5 5 1 39 7 1 1 2 1 1 8 8 8	Honolulu, Hawaii Long Beach, Calif. Los Angeles, Calif. Paradetras, Calif. Portland, Oreg. Sacramento, Calif. San Diego, Calif. San Francisco, Calif. Sants Cruz, Calif. Seattle, Wash. Tiscoma, Wash. Tiscoma, Wash.	102 U U 125 108 187	64 71 1 1 78 86 86 117 100 163 23 80 90 90 90	22 1 15 1 U 2 30 3 12 7 38 7 50 3 37 2 16 8 10	4 9 U U 13 8 277 200 199 - 4 1 5	4 3 U U 2 1 5 1 1 1 1 1 5 3 18	5 4 U U 1 1 2 2 4 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2	

*Mortality data in this table are voluntarily reported from 121 cities in the United States, most of which have populations of 100,000 or more. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included.

*Presumonia and influenza.

*Because of changes in reporting methods in these 3 Pennsylvania cities, these numbers are partial counts for the current week. Complete counts will be available in 4 to 8 weeks.

*Total includes unknown ages.

*U: Unaveilable -: no reported cases

Kombucha Tea — Continued

Prevalence of Kombucha Tea Drinking

To assess the prevalence of Kombucha tea drinking in the town, a 1% sample of households (n=129) was contacted by telephone using random-digit dialing. The mean age of the respondents was 51.2 years (standard deviation=±19.5 years), and 91 (70%) were women. Five persons (3.8%; 95% confidence interval [CI]=1.4%—8.4%) reported that at least one household member had tried Kombucha tea. Of these, two (1.6% of total sample; 95% CI=0.3%—5.0%) were persons who had regularly consumed the tea. Both had stopped drinking the tea after <2 weeks—one because of the tea's taste and one because of symptoms unrelated to those of the two patients.

Reported by: RW Currier, DVM, J Goddard, K Buechler, MP Quinlisk, MD, State Epidemiologist, lowa Dept of Public Health; SL Wolfe, MD, Spencer; TJ Carroll, MD, T Bennett, MD, Office of the Medical Examiner, Sioux City; J Stokes, MD, Univ of Iowa, Iowa City. Center for Food Safety, Food and Drug Administration. Health Studies Br, Div of Environmental Hezards and Health Effects, National Center for Environmental Health, CDC.

Editorial Note: The Kombucha "mushroom" is a symbiotic colony of several species of yeast and bacteria that are bound together by a surrounding thin membrane. Although the composition of the Kombucha colony varies, some of the species reportedly found in the mushroom include S. ludwigii, S. pombe, Bacterium xylinum, B. gluconicum, B. xylinoides, B. katogenum, Pichia fermentans, and Torula sp. (1). Kombucha tea can contain up to 1.5% alcohol and a variety of other metabolites (e.g., ethyl acetate, acetic acid, and lactate). During incubation, the thin, gelatinous mushroom floats in the tea and duplicates itself by producing a "baby" on top of the original mushroom. These offspring are then given to other persons for starting their own cultures. Although there are at least two commercial producers of Kombucha mushrooms in the United States, the sharing of the mushrooms is believed to have helped to promote its popularity in the United States.

Beneficial effects attributed to consumption of Kombucha tea have included prevention of cancer, relief of arthritis, treatment of insomnia, and stimulation of regrowth of hair (1-3). Because the tea is believed to stimulate the immune system, it has become popular among persons with human immunodeficiency virus infection (3). In addition, the investigation in lows suggests that the tea has become popular among the elderly (who are less likely to try alternative therapies) (4).

FDA has evaluated the practices of the commercial producers of the Kombucha mushroom and has found no pathogenic organisms or hygiene violations (5). However, because the tea is produced under varying conditions in individual homes, contamination with pathogenic organisms such as *Aspergillus* is possible. When prepared as directed, the pH of the tea decreases to 1.8 in 24 hours. Although this level of acidity should prevent the survival of most potentially contaminating organisms, tea drinkers have reported molds growing on the Kombucha (CDC, unpublished data).

Because folk medicines and herbal remedies, including Kombucha tea, are considered neither a food nor a drug (6–8), they are not routinely evaluated by FDA or the U.S. Department of Agriculture. Although the investigation described in this report did not establish a causal link between the illness of the two women and their consumption of Kombucha tea, reasons for the occurrence and severity of the lactic acidosis in both cases have not been determined. Drinking this tea in quantities typically consumed (approximately 4 oz daily) may not cause adverse effects in healthy persons; however, the potential health risks are unknown for those with preexisting health problems or those who drink excessive quantities of the tea.

Kombucha Tea — Continued

Because of the acidity of Kombucha tea, it should not be prepared or stored in containers made from materials such as ceramic or lead crystal, which both contain toxic elements than can leach into the tea. Because of the increasing use of this tea (even in groups that usually do not use alternative therapies), health-care professionals should consider consumption of Kombucha tea in the differential diagnosis of persons with unexplained lactic acidosis. Physicians and the public should report adverse health effects associated with consumption of Kombucha tea to FDA's MedWatch program, telephone (800) 332-1088 or (301) 738-7553.

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Update: Respiratory Syncytial Virus Activity — United States, 1995–96 Season

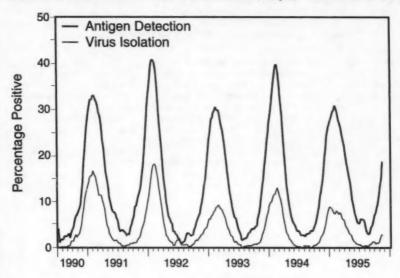
Respiratory syncytial virus (RSV), a common cause of winter outbreaks of acute respiratory disease, is associated each year with an estimated 90,000 hospitalizations and 4500 deaths from lower respiratory tract disease in both infants and young children in the United States (1). Outbreaks occur annually throughout the United States, and community activity usually peaks within 1 month of the national peak in January or February (Figure 1) (2). RSV activity in the United States is monitored by the National Respiratory and Enteric Virus Surveillance System (NREVSS), a voluntary, laboratory-based system. This report presents provisional surveillance results from the NREVSS for RSV during July 1–December 1, 1995, and summarizes trends in RSV from July 1990 through June 1995.

Since July 1, 1990, a total of 107 hospital-based and public health laboratories in 47 states have participated in the NREVSS and have reported weekly to CDC the number of specimens tested for RSV by the antigen-detection and virus-isolation methods and the number of positive results. Widespread RSV activity is defined by the NREVSS as the first of 2 consecutive weeks when at least half of participating laboratories report any RSV detections. This definition generally indicates a mean percentage of specimens positive by antigen detection >10%.

During the previous five seasons (i.e., July 1990–June 1995), onset of widespread RSV activity began in November and continued a mean of 22 weeks, until April or early May (Figure 1). Activity peaked each year from late January through mid-February. For the current reporting period (July 1–December 1, 1995), 72 laboratories

Respiratory Syncytial Virus — Continued

FIGURE 1. Percentage* of specimens positive for respiratory syncytial virus, by method of confirmation and month — United States, July 1, 1990–December 1, 1995



^{*}Laboratory-group mean, smoothed using a 5-week moving average.

in 44 states reported results of testing for RSV. Since October 21, more than half of the participating laboratories reported detections of RSV on a weekly basis, indicating the onset of RSV activity for the 1995–96 season.

Reported by: National Respiratory and Enteric Virus Surveillance System collaborating laboratories. Respiratory and Enteric Viruses Br, Div of Viral and Rickettsial Diseases, National Center for Infectious Diseases, CDC.

Editorial Note: During the RSV season, health-care providers should consider RSV in the differential diagnosis of acute respiratory disease in both children and adults. Most severe manifestations of RSV infection (e.g., pneumonia and bronchiolitis) occur in infants aged 2–6 months; however, children of any age with underlying cardiac or pulmonary disease or who are immunocompromised are at risk for serious complications from this infection. Because natural infection with RSV provides limited protective immunity, RSV may cause repeated symptomatic infections. In adults, RSV usually causes upper respiratory tract manifestations but may cause lower respiratory tract disease. Infection in immunocompromised persons can be associated with high death rates.

RSV is a common, but preventable, cause of nosocomially acquired infection; the risk for nosocomial transmission is increased during community outbreaks. Sources for nosocomially acquired infection include infected patients, staff, visitors, or contaminated fomites. Nosocomial outbreaks or transmission of RSV can be controlled

Respiratory Syncytial Virus - Continued

with strict attention to contact-isolation procedures (3). In addition, chemotherapy with ribavirin may be indicated for some patients (e.g., those at high risk for severe complications or who are seriously ill with this infection) (4). Prophylaxis with intravenous RSV immunoglobulin for high-risk patients may become available during future RSV seasons (5), and vaccines for RSV are being developed (6).

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Monthly Immunization Table

To track progress toward achieving the goals of the Childhood Immunization Initiative (CII), CDC publishes monthly a tabular summary of the number of cases of all diseases preventable by routine childhood vaccination reported during the previous month and year-to-date (provisional data). In addition, the table compares provisional data with final data for the previous year and highlights the number of reported cases among children aged <5 years, who are the primary focus of CII. Data in the table are reported through the National Electronic Telecommunications System for Surveillance (NETSS).

Number of reported cases of diseases preventable by routine childhood vaccination — United States, October 1994 and 1994–1995*

	No. cases, October	Total January-	cases October	No. cases among children aged <5 years ¹ January-October			
Disease	1995	1994	1995	1994	1995		
Congenital rubella							
syndrome	2	3	6	3	5		
Diphtheria	0	2	0	1	0		
Haemophilus influenzae§	86	939	974	247	233		
Hepatitis B¶	780	9423	8132	99	60		
Measles	11	875	280	207	101		
Mumps	63	1203	685	187	130		
Pertussis	438	3363	3398	1813	1951		
Poliomyelitis, paralytic**	0	1	0	0	0		
Rubella	5	209	135	24	17		
Tetanus	3	35	26	0	2		

^{*}Data for 1994 and 1995 are provisional.

[†]For 1994 and 1995, age data were available for ≥93% cases.

⁵Invasive disease; *H. influenzae* serotype is not routinely reported to the National Notifiable Diseases Surveillance System. Of 233 cases among children aged <5 years, serotype was reported for 56 cases, and of those, 33 were type b, the only serotype of *H. influenzae* preventable by vaccination.

Because most hepatitis B virus infections among infants and children aged <5 years are asymptomatic (although likely to become chronic), acute disease surveillance does not reflect the incidence of this problem in this age group or the effectiveness of hepatitis B vaccination in infants.

^{**}One case with onset in 1994 has been confirmed; this case was vaccine-associated. An additional six suspected cases are under investigation. In 1993, three of 10 suspected cases were confirmed; two of the confirmed cases were vaccine-associated, and one was imported. The imported case occurred in a 2-year-old Nigerian child brought to the United States for care of his paralytic illness; no poliovirus was isolated from the child.

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